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clinical procedures, minor surgical procedures not requiring muscle relaxation, and minor dental procedures not requiring intubation. The intravenous route of administration is more efficacious for dental care.

(3) *Limitations*. Federal law restricts this drug to use by or on the order of a licensed veterinarian.

[61 FR 21075, May 9, 1996, as amended at 79 FR 16191, Mar. 25, 2014]

§ 522.1350 Melatonin implant.

- (a) *Specifications*. The drug is a silicone rubber elastomer implant containing 2.7 milligrams of melatonin.
- (b) *Sponsor*. See No. 053923 in §510.600(c) of this chapter.
- (c) Conditions of use—(1) Amount. One implant per mink.
- (2) *Indications for use.* For use in healthy male and female kit and adult female mink (*Mustela vison*) to accelerate the fur priming cycle.
- (3) *Limitations*. For subcutaneous implantation in mink only. Do not implant potential breeding stock. Do not use in food-producing animals.

[59 FR 37422, July 22, 1994]

§ 522.1362 Melarsomine powder for injection.

- (a) Specifications. The drug consists of a vial of lyophilized powder containing 50 milligrams of melarsomine dihydrochloride which is reconstituted with the provided 2 milliliters of sterile water for injection.
- (b) Sponsor. See No. 050604 in §510.600(c) of this chapter.
- (c) Conditions of use—(1) Amount. Administer only by deep intramuscular injection in the lumbar muscles (L_{3} – L_{5}).
- (2) Indications. Treatment of stabilized, class 1, 2, and 3 heartworm disease (asymptomatic to mild, moderate, and severe, respectively) caused by immature (4 month-old, stage L₅) to mature adult infections of Dirofilaria immits in dogs.
- (3) *Limitations*. Federal law restricts this drug to use by or on the order of a licensed veterinarian.

[60 FR 49340, Sept. 25, 1995, as amended at 79 FR 16191, Mar. 25, 2014]

§ 522.1367 Meloxicam.

- (a) Specifications. Each milliliter of solution contains 5.0 milligrams (mg) meloxicam.
- (b) *Sponsor*. See Nos. 000010, 016729, and 055529 in §510.600(c) of this chapter.
- (c) Conditions of use—(1) Dogs—(i) Amount. Administer 0.09 mg per pound (mg/lb) body weight (0.2 mg per kilogram (mg/kg)) by intravenous or subcutaneous injection on the first day of treatment. For treatment after day 1, administer meloxicam suspension orally at 0.045 mg/lb (0.1 mg/kg) body weight once daily as in §520.1350(c) of this chapter.
- (ii) *Indications for use*. For the control of pain and inflammation associated with osteoarthritis.
- (iii) *Limitations*. Federal law restricts this drug to use by or on the order of a licensed veterinarian.
- (2) Cats—(i) Amount. Administer 0.14 mg/lb (0.3 mg/kg) body weight as a single, one-time subcutaneous injection.
- (ii) Indications for use. For the control of postoperative pain and inflammation associated with orthopedic surgery, ovariohysterectomy, and castration when administered prior to surgery.
- (iii) *Limitations*. Federal law restricts this drug to use by or on the order of a licensed veterinarian.

[68 FR 68724, Dec. 10, 2003, as amended at 69 FR 69523, Nov. 30, 2004; 78 FR 5715, Jan. 28, 2013]

§522.1372 Mepivacaine.

- (a) Specifications. Each milliliter (mL) of solution contains 20 milligrams mepivacaine hydrochloride.
- (b) Sponsor. See No. 054771 in \$510.600(c) of this chapter.
- (c) Conditions of use in horses—(1) Amount. For nerve block, 3 to 5 mL; for epidural anesthesia, 5 to 20 mL; for intra-articular anesthesia, 10 to 15 mL; for infiltration, as required; for anesthesia of the laryngeal mucosa prior to ventriculectomy, by topical spray, 25 to 40 mL, by infiltration, 20 to 50 mL.
- (2) Indications for use. For use as a local anesthetic for infiltration, nerve block, intra-articular and epidural anesthesia, and topical and/or infiltration anesthesia of the laryngeal mucosa prior to ventriculectomy.

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(3) Limitations. Not for use in horses intended for human consumption. Federal law restricts this drug to use by or on the order of a licensed veterinarian.

[71 FR 39547, July 13, 2006, as amended at 79 FR 16191, Mar. 25, 2014]

§522.1380 Methocarbamol.

- (a) Specifications. Each milliliter of solution contains 100 milligrams (mg) of methocarbamol.
- (b) Sponsor. See No. 054771 in \$510.600(c) of this chapter.
- (c) Conditions of use—(1) Amount—(1) Dogs and cats. Administer by intravenous injection 20 mg per pound of body weight for moderate conditions or 25 to 100 mg per pound of body weight for severe conditions (tetanus and strychnine poisoning). The total cumulative dose should not to exceed 150 mg per pound of body weight.
- (ii) Horses. Administer by intravenous injection 2 to 10 mg per pound of body weight for moderate conditions or 10 to 25 mg per pound of body weight for severe conditions (tetanus). Additional amounts may be needed to relieve residual effects and to prevent recurrence of symptoms.
- (2) Indications for use. As an adjunct for treating acute inflammatory and traumatic conditions of the skeletal muscles and to reduce muscular spasms.
- (3) *Limitations*. Do not use in horses intended for human consumption. Federal law restricts this drug to use by or on the order of a licensed veterinarian.

 $[79\;\mathrm{FR}\;16191,\,\mathrm{Mar}.\;25,\,2014]$

§ 522.1410 Methylprednisolone.

- (a) Specifications. Each milliliter of suspension contains 20 or 40 milligrams (mg) of methylprednisolone acetate.
- (b) Sponsors. See Nos. 054628 and 054771 in $\S 510.600(c)$ of this chapter.
 - (c) [Reserved]
- (d) Conditions of use—(1) Dogs—(i) Amount. Administer 2 to 40 mg (up to 120 mg in extremely large breeds or dogs with severe involvement) by intramuscular injection or up to 20 mg by intrasynovial injection.
- (ii) Indications for use. For treatment of inflammation and related disorders; treatment of allergic and dermatologic disorders; and as supportive therapy to

- antibacterial treatment of severe infections.
- (iii) *Limitations*. Federal law restricts this drug to use by or on the order of a licensed veterinarian.
- (2) *Cats*—(i) *Amount*. Administer 10 to 20 mg by intramuscular injection.
- (ii) Indications for use. For treatment of inflammation and related disorders; treatment of allergic and dermatologic disorders; and as supportive therapy to antibacterial treatment of severe infections.
- (iii) *Limitations*. Federal law restricts this drug to use by or on the order of a licensed veterinarian.
- (3) *Horses*—(i) *Amount*. Administer 200 mg by intramuscular injection or 40 to 240 mg by intrasynovial injection.
- (ii) *Indications for use*. For treatment of inflammation and related disorders.
- (iii) *Limitations*. Do not use in horses intended for human consumption. Federal law restricts this drug to use by or on the order of a licensed veterinarian.
- [43 FR 59058, Dec. 19, 1978, as amended at 51 FR 741, Jan. 8, 1986; 53 FR 40728, Oct. 18, 1988; 62 FR 35076, June 30, 1997; 76 FR 53051, Aug. 25, 2011; 78 FR 21060, Apr. 9, 2013; 79 FR 16191, Mar. 25, 2014]

§ 522.1450 Moxidectin solution.

- (a) Specifications. Each milliliter of solution contains 10 milligrams (mg) moxidectin.
- (b) *Sponsor*. See No. 000010 in §510.600(c) of this chapter.
- (c) Related tolerances. See §556.426 of this chapter.
- (d) Special considerations. See §500.25 of this chapter.
- (e) Conditions of use in beef and non-lactating dairy cattle— (1) Amount. Administer 0.2 mg/kg of body weight (0.2 mg/2.2 pound) as a single, subcutaneous injection.
- (2) Indications for use. For treatment and control of gastrointestinal Ostertagiaroundworms: ostertaai (adults, fourth-stage larvae, and inhib-Haemonchus ited larvae). placei (adults), Trichostrongylus axei (adults and fourth-stage larvae). Trichostrongulus colubriformis (adults fourth-stage larvae), Cooperia oncophora (adults), Cooperia pectinata (adults), Cooperia punctata (adults and fourth-stage larvae), Cooperia spatulata (adults), Cooperia surnabada (adults and